AMENDMENTS TO THE CLAIMS

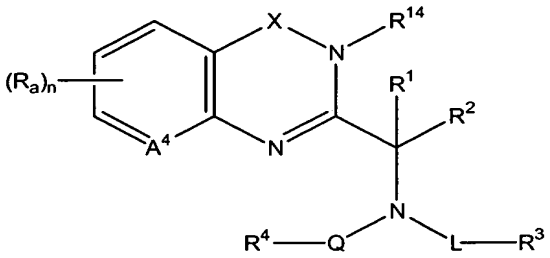
Please cancel claims 156-159, 162-170, 173-190, 193-197, and 202 without prejudice.

Please amend claims 136, 138, 139, and 154 as shown below.

Please add new claims 205-226 as shown in the following list of claims:

1.-135. (Canceled).

136. (Currently Amended) A compound having the formula:



or a pharmaceutically acceptable salt thereof wherein:

A^4 is N;

X is -C(O)- or $-CH_2-$;

R^1 and R^2 are members independently selected from the group consisting of H and (C_1-C_4)alkyl;

R^3 is a member selected from the group consisting of hydroxy, (C_1-C_8)alkoxy, amino, (C_1-C_8)alkylamino, di(C_1-C_8)alkylamino, (C_2-C_8)heteroalkyl, (C_3-C_9)heterocyclyl, (C_1-C_8)acylamino, amidino, guanidino, ureido, cyano, heteroaryl, $-CONR^9R^{10}$ and $-CO_2R^{11}$;

R^4 is a member selected from the group consisting of (C_1-C_{20})alkyl, (C_2-C_{20})heteroalkyl, heteroaryl, aryl, heteroaryl(C_1-C_6)alkyl, heteroaryl(C_2-C_6)heteroalkyl, aryl(C_1-C_6)alkyl and aryl(C_2-C_6)heteroalkyl;

each R^9, R^{10} and R^{11} is independently selected from the group consisting of H, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, heteroaryl, aryl, heteroaryl(C_1-C_6)alkyl, heteroaryl(C_2-C_8)heteroalkyl, aryl(C_1-C_8)alkyl and aryl(C_2-C_8)heteroalkyl;

R^{14} is a substituted or unsubstituted ~~aryl or heteroaryl~~ member selected from the group consisting of phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl;

Q is -C(O)-;

L is (C_1-C_8)alkylene;

the subscript n is an integer from 0 to 4; and

each R_a is independently selected from the group consisting of halogen, -OR', -OC(O)R', -NR'R", -SR', -R', -CN, -NO_2, -CO_2R', -CONR'R", -C(O)R', -OC(O)NR'R", -NR"C(O)R', -NR"$C(O)_2$R', ,-NR'-C(O)NR"R"', -NH-$C(NH_2)$=NH, -NR'$C(NH_2)$=NH, -NH-$C(NH_2)$=NR', -S(O)R', -$S(O)_2$R', -$S(O)_2$NR'R", -N_3, -$CH(Ph)_2$, perfluoro(C_1-C_4)alkoxy and perfluoro(C_1-C_4)alkyl, wherein R', R" and R"' are each independently selected from the group consisting of H, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C_1-C_4)alkyl and (unsubstituted aryl)oxy-(C_1-C_4)alkyl.

137. (Previously Added) The compound of Claim 136, wherein X is –C(O)-.

138. (Currently Amended) The compound of Claim 136, wherein R^{14} is a substituted or unsubstituted phenyl. ~~member selected from the group consisting of phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl.~~

139. (Currently Amended) The compound of Claim 137, wherein R^{14} is a substituted or unsubstituted phenyl. ~~member selected from the group consisting of phenyl, pyridyl, thiazolyl, thienyl and pyrimidinyl.~~

140. (Previously Added) The compound of Claim 136, wherein R^3 is (C_1-C_8)acylamino.

141. (Previously Added) The compound of Claim 136, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo(C_1-C_4)alkyl, halo(C_1-C_4)alkoxy, cyano, nitro and phenyl.

142. (Previously Added) The compound of Claim 136, wherein R^{14} is selected from the group consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl and substituted thienyl, wherein the substituents are selected from the group consisting of cyano, halogen, (C_1-C_8)alkoxy, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

143. (Previously Added) The compound of Claim 136, wherein R^{14} is substituted phenyl, wherein the substituents are selected from the group consisting of cyano, halogen, (C_1-C_8)alkoxy, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

144. (Previously Added) The compound of Claim 136, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo(C_1-C_4)alkyl, halo(C_1-C_4)alkoxy, cyano, nitro and phenyl, and R^{14} is substituted phenyl, wherein the substituents are selected from the group consisting of cyano, halogen, (C_1-C_8)alkoxy, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

145. (Previously Added) The compound of Claim 136, wherein R^1 is selected from the group consisting of methyl, ethyl and propyl, and R^2 is hydrogen.

146. (Previously Added) The compound of Claim 136, wherein R^1 and R^2 are each methyl.

147. (Previously Added) The compound of Claim 136, wherein L is (C_1-C_4)alkylene.

148. (Previously Added) The compound of Claim 136, wherein R^3 is a member selected from the group consisting of (C_1-C_8)alkoxy, (C_3-C_9)heterocyclyl and heteroaryl.

149. (Previously Added) The compound of Claim 136, wherein R^3 is heteroaryl.

150. (Previously Added) The compound of Claim 136, wherein R^3 is heteroaryl and R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo(C_1-C_4)alkyl, halo(C_1-C_4)alkoxy, cyano, nitro and phenyl.

151. (Previously Added) The compound of Claim 136, wherein R^3 is selected from the group consisting of substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl.

152. (Previously Added) The compound of Claim 136, wherein R^1 and R^2 are each independently selected from the group consisting of H, methyl and ethyl; R^{14} is phenyl; L is methylene, ethylene or propylene; R^3 is selected from the group consisting of substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; and R^4 is substituted or

unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo(C_1-C_4)alkyl, halo(C_1-C_4)alkoxy, cyano, nitro and phenyl.

153. (Previously Added) A pharmaceutical composition comprising the compound of Claim 136 and a pharmaceutically acceptable carrier or diluent.

154. (Currently Amended) A method of treating psoriasis, rheumatoid arthritis, inflammatory bowel disease, asthma, organ transplant conditions, or multiple sclerosis ~~an inflammatory or immune condition or disease~~ in a subject, said method comprising administering to a subject in need of such treatment a therapeutically effective amount of the compound of Claim 136.

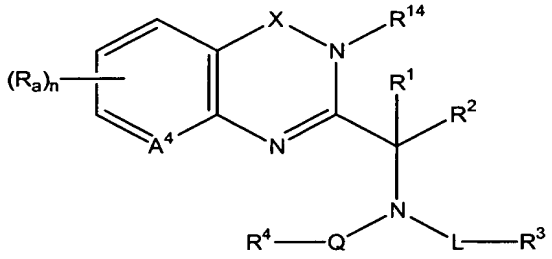
155. (Previously Added) The method of Claim 154, wherein said compound is administered orally, parenterally or topically.

156.-202. (Canceled).

203. (Previously Added) A method for the modulation of CXCR3 function in a cell, comprising contacting said cell with a compound of Claim 136.

204. (Previously Added) A method for the modulation of CXCR3 function, comprising contacting a CXCR3 protein with a compound of Claim 136.

205. (New) A compound having the formula:



or a pharmaceutically acceptable salt thereof wherein:

A^4 is N;

X is -C(O)- or $-CH_2-$;

R^1 and R^2 are members independently selected from the group consisting of H and (C_1-C_4)alkyl;

R^3 is a member selected from the group consisting of hydroxy, (C_1-C_8)alkoxy, amino, (C_1-C_8)alkylamino, di(C_1-C_8)alkylamino, (C_2-C_8)heteroalkyl, (C_3-C_9)heterocyclyl, (C_1-C_8)acylamino, amidino, guanidino, ureido, cyano, heteroaryl, $-CONR^9R^{10}$ and $-CO_2R^{11}$;

R^4 is a member selected from the group consisting of (C_1-C_{20})alkyl, (C_2-C_{20})heteroalkyl, heteroaryl, aryl, heteroaryl(C_1-C_6)alkyl, heteroaryl(C_2-C_6)heteroalkyl, aryl(C_1-C_6)alkyl and aryl(C_2-C_6)heteroalkyl;

each R^9, R^{10} and R^{11} is independently selected from the group consisting of H, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, heteroaryl, aryl, heteroaryl(C_1-C_6)alkyl, heteroaryl(C_2-C_8)heteroalkyl, aryl(C_1-C_8)alkyl and aryl(C_2-C_8)heteroalkyl;

R^{14} is substituted or unsubstituted aryl or heteroaryl;

Q is -C(O)-;

L is (C_1-C_8)alkylene;

the subscript n is an integer from 0 to 4; and

each R_a is independently selected from the group consisting of halogen, -OR', -OC(O)R', -NR'R", -SR', -R', -CN, $-NO_2$, $-CO_2R'$, -CONR'R", -C(O)R', -OC(O)NR'R", -NR"C(O)R', $-NR"C(O)_2R'$, ,-NR'-C(O)NR"R"', $-NH-C(NH_2)=NH$, $-NR'C(NH_2)=NH$, $-NH-C(NH_2)=NR'$, -S(O)R', $-S(O)_2R'$, $-S(O)_2NR'R"$, $-N_3$, $-CH(Ph)_2$, perfluoro(C_1-C_4)alkoxy and perfluoro(C_1-C_4)alkyl, wherein R', R" and R"' are each independently selected from the group consisting of H, (C_1-C_8)alkyl, (C_2-C_8)heteroalkyl, unsubstituted aryl, unsubstituted heteroaryl, (unsubstituted aryl)-(C_1-C_4)alkyl and (unsubstituted aryl)oxy-(C_1-C_4)alkyl.

206. (New) The compound of Claim 205, wherein X is -C(O)-.

207. (New) The pharmaceutical composition of Claim 153, wherein X is -C(O)-.

208. (New) The pharmaceutical composition of Claim 153, wherein R^{14} is a substituted or unsubstituted phenyl.

209. (New) The pharmaceutical composition of Claim 153, wherein R^3 is (C_1-C_8)acylamino.

210. (New) The pharmaceutical composition of Claim 153, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo(C_1-C_4)alkyl, halo(C_1-C_4)alkoxy, cyano, nitro and phenyl.

211. (New) The pharmaceutical composition of Claim 153, wherein R^{14} is selected from the group consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl and substituted thienyl, wherein the substituents are selected from the group consisting of cyano, halogen, $(C_1\text{-}C_8)$alkoxy, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

212. (New) The pharmaceutical composition of Claim 153, wherein R^{14} is substituted phenyl, wherein the substituents are selected from the group consisting of cyano, halogen, $(C_1\text{-}C_8)$alkoxy, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

213. (New) The pharmaceutical composition of Claim 153, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo$(C_1\text{-}C_4)$alkyl, halo$(C_1\text{-}C_4)$alkoxy, cyano, nitro and phenyl, and R^{14} is substituted phenyl, wherein the substituents are selected from the group consisting of cyano, halogen, $(C_1\text{-}C_8)$alkoxy, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

214. (New) The pharmaceutical composition of Claim 153, wherein R^1 is selected from the group consisting of methyl, ethyl and propyl, and R^2 is hydrogen.

215. (New) The pharmaceutical composition of Claim 153, wherein L is $(C_1\text{-}C_4)$alkylene.

216. (New) The pharmaceutical composition of Claim 153, wherein X is -C(O)-; R^1 and R^2 are each independently selected from the group consisting of H, methyl and ethyl; R^{14} is selected from the group consisting of substituted and unsubstituted phenyl; L is methylene, ethylene or propylene; R^3 is selected from the group consisting of substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; and R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo$(C_1\text{-}C_4)$alkyl, halo$(C_1\text{-}C_4)$alkoxy, cyano, nitro and phenyl.

217. (New) The method of Claim 154, wherein X is -C(O)-.

218. (New) The method of Claim 154, wherein R^{14} is a substituted or unsubstituted phenyl.

219. (New) The method of Claim 154, wherein R^3 is $(C_1\text{-}C_8)$acylamino.

220. (New) The method of Claim 154, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo$(C_1\text{-}C_4)$alkyl, halo$(C_1\text{-}C_4)$alkoxy, cyano, nitro and phenyl.

221. (New) The method of Claim 154, wherein R^{14} is selected from the group consisting of substituted phenyl, substituted pyridyl, substituted thiazolyl and substituted thienyl, wherein the substituents are selected from the group consisting of cyano, halogen, $(C_1\text{-}C_8)$alkoxy, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

223. (New) The method of Claim 154, wherein R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo$(C_1\text{-}C_4)$alkyl, halo$(C_1\text{-}C_4)$alkoxy, cyano, nitro and phenyl, and R^{14} is substituted phenyl, wherein the substituents are selected from the group consisting of cyano, halogen, $(C_1\text{-}C_8)$alkoxy, $(C_1\text{-}C_8)$alkyl, $(C_2\text{-}C_8)$heteroalkyl, $CONH_2$, methylenedioxy and ethylenedioxy.

224. (New) The method of Claim 154, wherein R^1 is selected from the group consisting of methyl, ethyl and propyl, and R^2 is hydrogen.

225. (New) The method of Claim 154, wherein L is $(C_1\text{-}C_4)$alkylene.

226. (New) The method of Claim 154, wherein X is -C(O)-; R^1 and R^2 are each independently selected from the group consisting of H, methyl and ethyl; R^{14} is selected from the group consisting of substituted and unsubstituted phenyl; L is methylene, ethylene or propylene; R^3 is selected from the group consisting of substituted or unsubstituted pyridyl and substituted or unsubstituted imidazolyl; and R^4 is substituted or unsubstituted benzyl, wherein said substituents are selected from the group consisting of halogen, halo$(C_1\text{-}C_4)$alkyl, halo$(C_1\text{-}C_4)$alkoxy, cyano, nitro and phenyl.